

Mass

- Mechanism Local destruction of lung parenchyma
- Radiological sign Any localized opacity not completely bordered by fissures or pleura

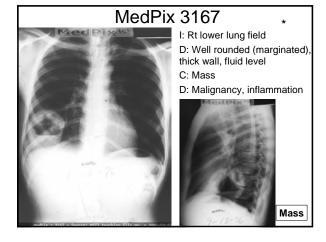


Mass

Mass Differential Diagnosis

- Malignancy Primary or secondary
- Granulomatous disease Infectious or noninfectious Active or inactive
- Other inflammation including pneumonia; abscess
- Benign neoplasm
- Congenital abnormality

Mass



Bronchgenic carcinoma

FINDINGS:

FINDINGS:
A cavitated round opacity is present at the right lung base. It overlies the back of the heart shadow on the lateral view. There are nodular opacities inside the cavity and an air-fluid level is also visible. The location is thus right lower lobe, with possible involvement of the posterior portion of the middle lobe.

PATTERN: The definition of a mass is satisfied.

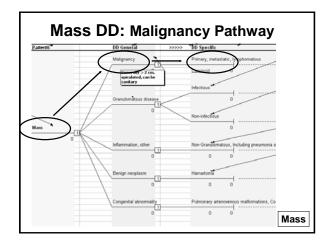
DIFFERENTIAL DIAGNOSIS:

Malignancy is favored over inflammation because of the irregularity of the inner wall of the cavity. The air-fluid level is not useful in differential diagnosis; it indicates only that the bronchus connected to the mass is either partially or intermittently obstructed.

DIAGNOSIS:

Bronchogenic carcinoma, adenomatous

Mass

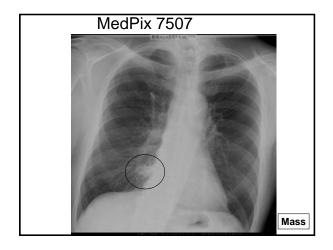


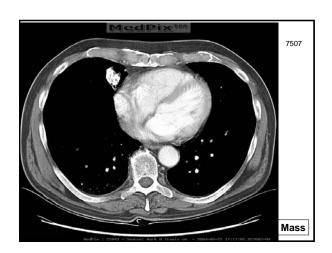
Malignancy: histological types

- Carcinoma (adenocarcinoma) (33%)
 - Bronchoalveolar cell carcinoma (BAC)
- Squamous cell (epidermoid) (30%)
- Large Cell carcinomas (4%)
 - usually classified as non-small cell types
- Small Cell carcinoma (13%)

<u>Thompson WH.</u> Respir Care. 2004 Nov;49(11):1349-53. Bronchioloalveolar carcinoma masquerading as pneumonia. <u>BAC</u>

http://www.archbronconeumol.org/cgi-bin/wdbcgi.exe/abn/abneng.mrevista.fulltext?pident=13076438





Mass Considerations: Calcifications

- Crucial appearance characteristics for inactivity
 - -Calcification
 - Central, lamellar
- Evolution
 - 2 year stability or regression

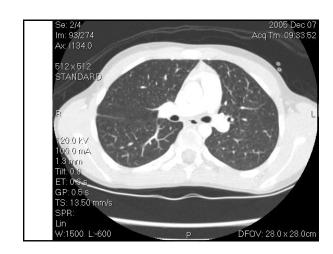
Pulmonary Hamartoma

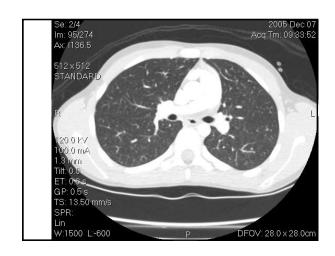
- Hamartomas are benign neoplasms
 90% found in lung
- · 5% of all solitary lung nodules.
- CR demonstrate well-circumscribed peripheral rounded or lobulated tumor.
- Frequently contain cartilage with fibrous connective tissue and various amounts of fat, smooth muscle, and seromucous glands.
- Approximately 30% contain calcium usually of the "popcorn" variety.
- · Seen most commonly in 4th and 5th decades of life.
- They are rare in children.

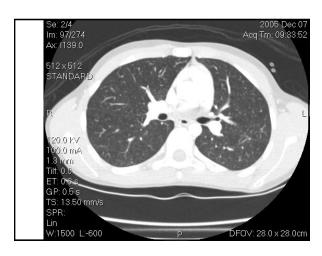
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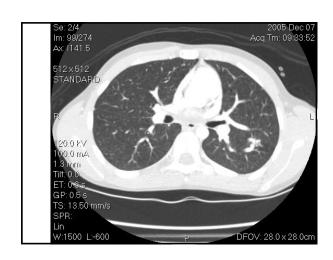
Mass

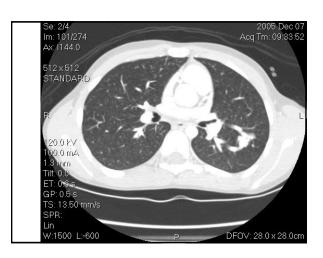


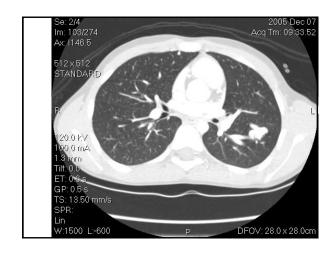


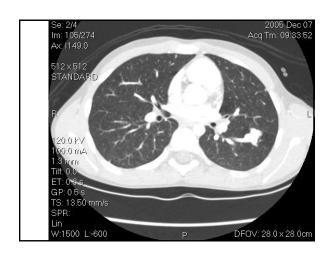






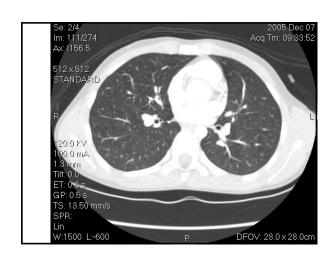






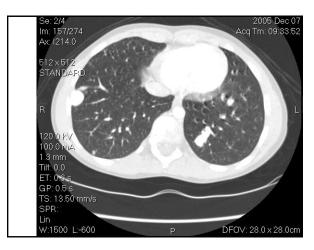














Pulmonary Arteriovenous Malformation

- Pulmonary AVM's are abnormal connections between the pulmonary arteries and veins.
- They are single in 65%, multiple in 35%.
 Twice as common in women than men, the majority are Congenital and are found in the lower lobes.
- Significantly, nearly 70% are associated with Hereditary Hemorrhagic Telangiectasia (Rendu-Osler-Weber disease), an autosomal dominant condition involving multiple AVM's in the brain, lung, skin, and liver.

The CT scan in this patient is highly suggestive and limits the differential diagnosis to pulmonary AVM.

The DDX for pulmonary AVM's can further be subdivided to Hereditary Hemorrhagic Telangiectasia and hepatopulmonary syndrome as the two most common causes.

History and physical exam findings support pulmonary AVM's and more specifically HHT

Hereditary Hemorrhagic Telangiectasia

- Also called Rendu-Osler-Weber syndrome
- · Autosomal Dominant inheritance
- Signs/Symptoms:
 - Epistaxis
 - Hemoptysis
 - Dyspnea
 - Clubbing
 - Skin telangiectasias
 - Gastrointestinal bleeding
 - Bruits/murmurs
 - High output CHF

Mass Considerations The Topographic Map Clinical variables Symptoms and signs Age, History Hemoptysis Prior exams (CXR, CT) Coughing Location (in chest) Dyspnea Multiplicity? SOB Mass DD is included in some Risk factors vascular and nodular patterns -Smoking -Occupation, exposure Also works the other way around: hobbies Not always black and -Previous carcinoma -Concurrent disease Factors narrowing down the differential dx. Mass

Consolidative (Alveolar) Pattern

Con-SOLID-ative

Mechanism

- -Produced in pure form by ALVEOLAR FILLING of density greater than air
- -May be mimicked by alveolar collapse,
- as in airway obstruction
 - -Rarely, manifests w confluent interstitial thickening

Consolidative

Consolidative (alveolar) Pattern Radiological signs:

Fluffy, cloud-like, coalescent opacities

Can get sharp edges when limited by fissures or pleura

Complete air bronchograms

Distribution: lobar

Obliterates pulmonary vasculature

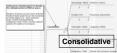
Differentiates from "ground glass" (not SOLID)

Consolidative

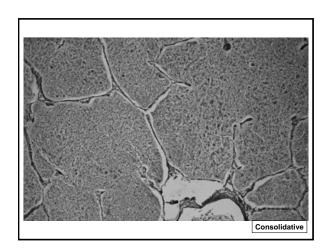
Consolidative (alveolar)

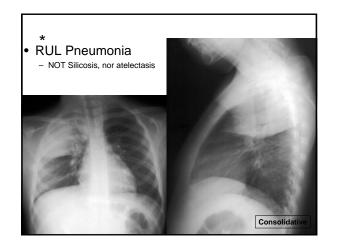
Pattern: Differential Diagnosis

- Hemorrhage **BLOOD** embolism, trauma
- Exudate PUS pneumonia, pneumonitis
- Transudate WATER congestion, ARDS
- Secretions **PROTEIN** Mucous plugging, Alveolar proteinosis
- Malignancy CELLS Bronchoalveolar cell carcinoma (BAC), Lymphoma









RUL Pneumonia

- Large area of opacification on the frontal view has both major and minor fissures as its inferior border.
- The lateral view demonstrates nicely the fissures of the right lung. Both RML and RLL remain well areated.
- MSU Top 10 CXR dx

 www.rad.msu.edu/.../ im_tutor/images/

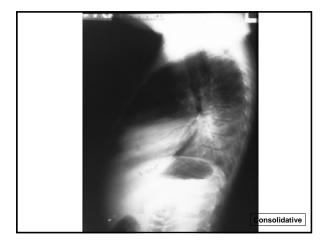
Consolidative

Bacterial pneumonia

- Streptococcus pneumoniae is the most common cause of bacterial pneumonia
- May present with mild to severe symptoms, including shaking chills, chattering teeth, severe chest pain, and a cough productive of rust-colored or greenish sputum
- May be febrile, diaphoretic, tachypneic, dyspneic, and/or cyanotic.

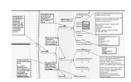
Consolidative



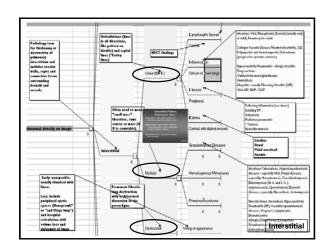


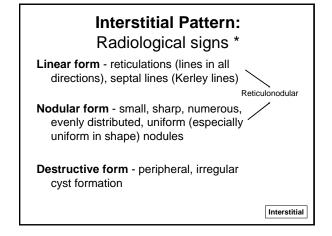
INTERSTITIAL PATTERN

- Mechanism:
 - -Thickening of lung interstices
 - Architectural destruction of interstitium
- Appearance: Lines, reticulations
 - -That are not vasculature; in addition of

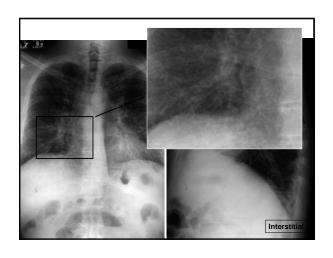


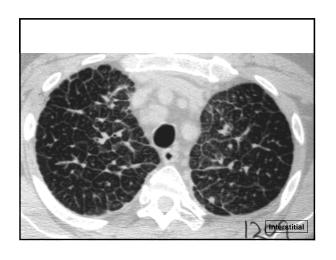
Interstitial

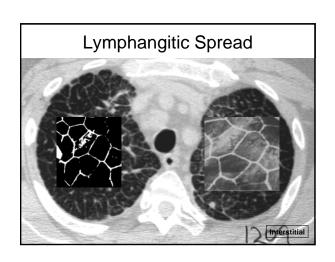


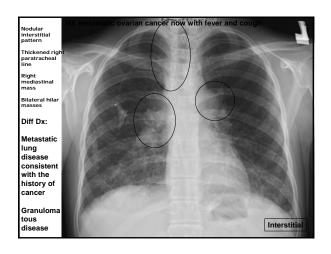


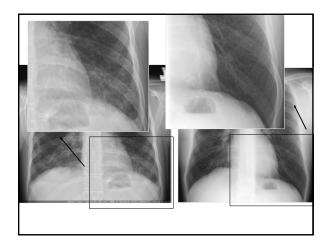


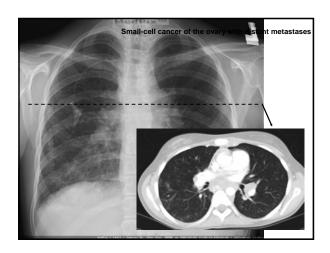


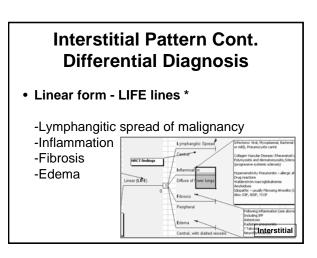


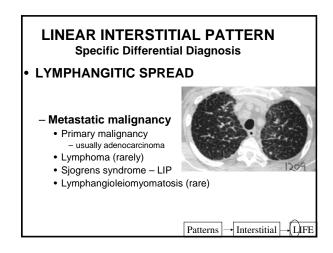


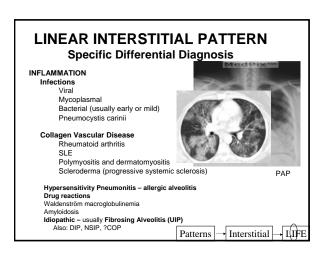












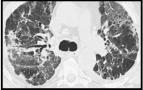


• FIBROSIS

Following inflammation **Including IPF Asbestosis** Radiation pneumonitis

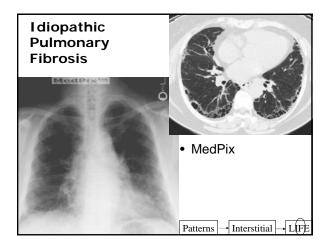
? Talcosis

Neurofibromatosis



Hypersensitivity Pneumonitis

→ Interstitial → LIFE Patterns |



LINEAR INTERSTITIAL PATTERN **Specific Differential Diagnosis**

EDEMA

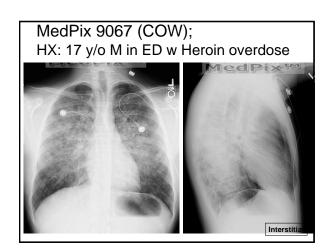
Mnemonic: NOT CARDIAC:

- Near drowning
 - Oxygen therapy
- Cardiac • Transfusion or Trauma (fat embolism)
- Renal

Anemia

- · Central nervous system disorder
- Allergic alveolitis Fluid overload
 - Renal failure
 - Drugs
 - Inhaled toxins
 - Aspiration or ARDS or Altitude sickness





Findings, DX

- · Bilateral patchy diffuse opacities predominantly in mid to upper lung fields
- DDG:
- • Noncardiogenic pulmonary edema
 - · Cardiogenic pulmonary edema
 - Allergic reaction
 - Lymphangitic spread
- DDS:
- Noncardiogenic Pulmonary Edema

http://rad.usuhs.mil/medpix/parent.php3?mode=cowpt&pt_id=9067&case=&recnum=0&imid=27311&showall=yes&hx=yes&dx=yes&th=1#diagnosis,

http://radiographics.rsnajnls.org/cgi/content/full/19/6/1507

Interstitial

Interstitial Pattern Cont. Differential Diagnosis

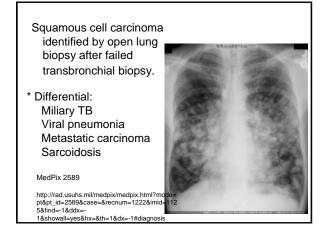
Nodular form

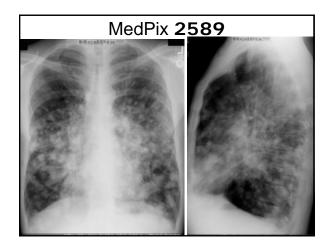
Granulomas

Hematogenous spread of malignancy Pneumonoconiosis

Since nodules are essentially small masses, include mass differential

Patterns - Interstitial - Nodular





INTERSTITIAL PATTERN: Nodular Specific Differential Diagnosis

Granulomatous Diseases Infectious

> Tuberculosis (miliary example below) Atypical mycobacterial diseases especially MAI



Patterns - Interstitial - Nodular

Miliary Nodular INTERSTITIAL PATTERN MedPix

TEMPEST

T TB, FUNGAL, VIRAL PNEUMONIAS

E EG

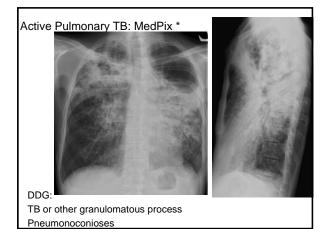
M METS (THYROID, RENAL)

P PNEUMOCONIOSES, PARASITES E EMBOLISM OF OILY CONTRAST (LIPOID PNEUMONIA)

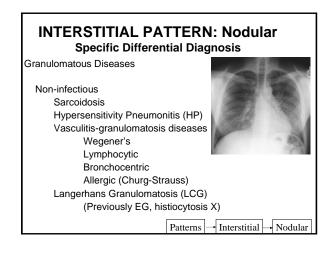
SARCOIDOSIS, SILICOSIS

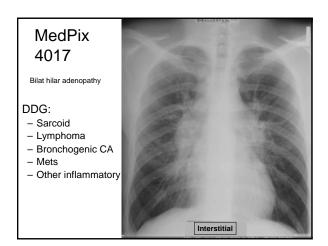
TUBEROUS SCLEROSIS

Interstitial



INTERSTITIAL PATTERN: Nodular Specific Differential Diagnosis Granulomatous Diseases Fungal diseases, especially: Histoplasmosis Coccidioidomycosis Blastomycosis (N. A. and S. A.) Cryptococcosis Sporotrichosis Bacterial diseases, especially: Nocardiosis Actinomycosis Patterns - Interstitial - Nodular

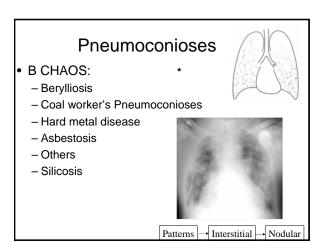




Sarcoid MedPix 4017

- A granulomatous disease of unclear etiology, most commonly recognized by its thoracic manifestations of interstitial lung disease and hilar and mediastinal adenopathy.
- A multisystem disease, with histologic evidence of sarcoid involvement of the liver and spleen seen in 50-80% of all surgical specimens, although most cases do not result in organ dysfunction.

Interstitial



Interstitial Pattern

Destructive form

Early appearance is nonspecific Late findings include peripheral cystic spaces

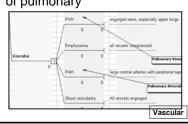
- Honeycomb or "End Stage Lung" with volume loss and deformity of lungs
- Chronic ILD: Low lung volumes
- · No airway wall thickening

Patterns Interstitial Destructive

VASCULAR PATTERN

 Mechanism - increased or decreased perfusion altering diameter (hence appearance) of pulmonary vessels

Larger diameter:
Big vessels
Smaller diameter:
Disappearance
More lucent
(Not displacement)



Vascular Pattern DDG: Examples

- Common examples
 - -Congestion engorged veins, especially upper lungs
 - -Emphysema diminished vessels
 - -Shunt vascularity all vessels enlarged
 - -Lymphangitic carcinoma irregular infiltration around vessels may resemble vessel enlargement

Vascular

Vascular Pattern DDG: Examples, cont.

Pulmonary Arterial hypertension - large central arteries with peripheral tapering

Thromboembolism - locally diminished vessels sometimes with large central vessels

Bronchial circulation - irregular vessels in unusual directions

Vascular

PAH vs. PVH

Pulmonary Arterial Hypertension

enlarged central and hilar vessels (best seen on lateral) enlarged middle mogul on PA pruned peripheral vessels (tapering to diminished) mosaic perfusion (CT) cor pulmonale

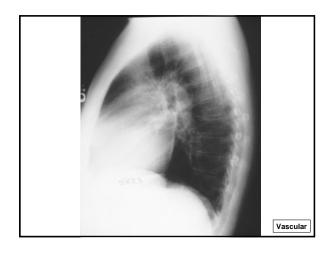
PA atherosclerosis

Pulmonary Venous Hypertension

septal lines smooth pleural thickening pleural effusion ground glass opacity dilated vessels, especially upper lungs hilar arteries not enlarged

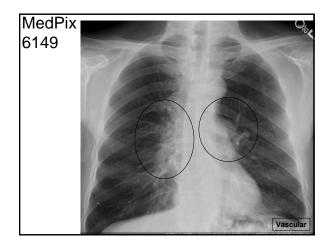
Vascular

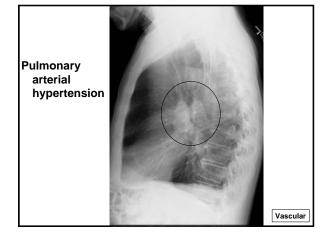












AIRWAY (BRONCHIAL) PATTERNS

- Mechanism:
- Complete or partial obstruction of airways
- Thickening of airway walls
 - or displacement of vessels due to overaeration, COPD,

Airway

Airway (bronchial) Pattern; Major Forms:

Complete airway obstruction

- opacity and decreased volume

Partial obstruction

- lucency and increased volume

Wall thickening

Show up as "tram tracks"
Bronchiolar (small airway) obstruction

Airway

Complete airway obstruction

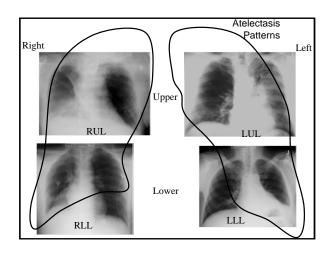
Opacities

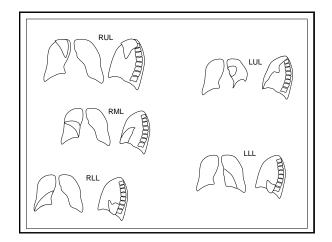
endobronchial malignancies, granulomas, inflammatory, benign or congenital masses, mucous plugs, foreign bodies

Decreased volume (atelectasis)

- can resemble mass or consolidation

Airway Complete



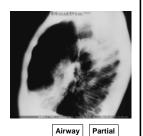


Partial airway obstruction

-Lucencies

- COPD, cysts, blebs, pneumatoceles





COPD with bullous emphysema

The lungs are hyperinflated and the diaphragms are markedly flattened, especially on the lateral view. There are numerous lucent "holes" in the lungs and the vessels are displaced and asymmetrical.

Air trapping is present, especially in multiple bullae with thin walls. These are the findings of bullous emphysema. Most such patients have COPD, the most common of all airway diseases.

DIFFERENTIAL DIAGNOSIS:

A few emphysematous patients have normal airways, with abnormal elasticity of alveolar walls, such as in alpha one antitrypsin deficiency.

DIAGNOSIS: COPD with bullous emphysema

Airway

http://rad.usuhs.mil/medpix/medpix.html?mode=single&recnum=1696&th=-1#top

DESTRUCTIVE AIRWAY DISEASE: BRONCHIECTASIS

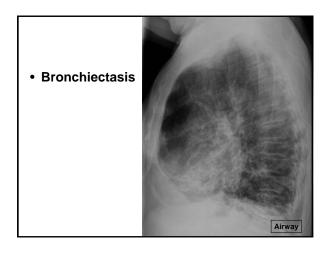
Thickening - bronchiectasis, chronic bronchitis

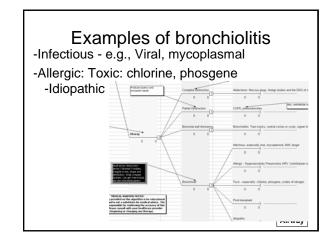
- · Central (perihilar) cystic spaces, increased lung volumes and thickened airway walls
 - circles and tram tracks
- · Distinct from destructive interstitial disease

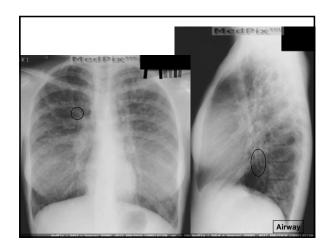
Note that air is the majority of the finding in airway pattern and density (bronchial walls) are the exception; as opposed to consolidation.

Airway Wall thickening





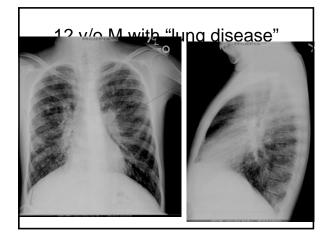




MedPix 9413 Cystic Fibrosis

- The radiographic findings are largely secondary to the bronchial obstructions by the thick adherent secretions. Pulmonary hyperinflation is evidenced in this case by the narrow heart and mediastinum and the slightly depressed diaphragm.
- The hila are prominent and there, is lobulated contour of the right hilum consistent with the presence of adenopathy which is common in these patients. The most prominent findings in this patient are the changes related to bronchiectasis.
- There is bronchial wall thickening seen as bronchial cuffing or "tram lines". Dilated bronchi are particularly well seen in the upper lungs where widened tubular and branching lucencies can be seen peripherally some of which containing tubular opacities representing impacted mucus.

Airway



Cystic Fibrosis - End-Stage

 Pt with end stage cystic fibrosis with pulmonary colonization with panresistant Pseudomonas cepacia.

Bronchiolar (small airway) obstruction

Airway ("alveolar") nodules: irregular in size, shape and distribution

• Small, irregular lucenies. Can get Tree-inbud, see also interstitial pattern

Airway Bronchiolar

Airway Patterns: CT

- -Thick-walled airways: "tram tracks" circular on end: "signet rings"
- -Cystic spaces centrally located
- -Cystic spaces with very thin walls or no apparent walls
- -Thin, stretched vessels

Airway

Summary

Presentation Format

Major CXR Patterns

Consolidative Interstitial Vascular **Airway**

Systematic Process, Methodical

Lung volumes Location/ distribution Patterns (type of opacity) DD General, then specific Definitions **Patterns** DDG

DDS

Cases, examples

Overview

References

http://rad.usuhs.mil/rad/handouts/ms-2_final.html

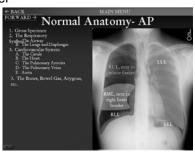
Case Studies: http://rad.usuhs.mil/rad/handouts/feigin/abnlcxr/myindex.htm

3 D anatomy:

http://vertex.biostr.washington.edu/cgi-bin/DA/imageform

Review Materials

- Chest Primer
- Anatomy
- MedPix

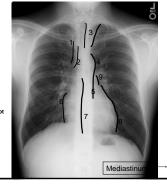


Review: Mediastinum Lines, Edges

Recommendation:

Test yourself before advancing to next slide

- SVC Edge Rt Paratracheal Line
- Lt Paratracheal Stripe (both red and black lines)
- Aortic Arch
 Descending Aorta
 (only left edge seen, and nalways)
- Rt Atrium
- Lt Ventricle
- Main Pulmonary Artery AKA: trunk, middle mogu



Mid